California Drug Recall Information

Recall Name
Hospira Recalls 0.9% Sodium Chloride Injection, USP 1000 mL, Flexible Container Due to Possible Brass Contamination

Recall Date	Product Description	Recalling Firm	Recall Reason
3/29/13	0.9% Sodium Chloride Injection, USP 1000 mL, Flexible Container **NDC # 0409-7983-09**	**Hospira, Inc.** Lake Forest, IL	*Confirmed customer report of several grey/brown particles identified as brass, was noted in solution*
Recall Class	**Product Identification**	**Distribution**	**Affected Dates**
n/a	Suspect Lot Recalled: • **25-037-JT (lot number may be followed by a "-01" or "-90")** • **Exp. date: January 1, 2015**	**CA**, nationwide	Distributed between January 2013 and March 2013

FOR ADDITIONAL INFORMATION, PLEASE VISIT:

http://www.fda.gov/Safety/Recalls/ucm345963.htm?source=govdelivery